

FIG. 1

Fok I site

↓
 nnnnnGGATGnnnnnnnnnnnnnnnnnnnnnnnnnnnn
 nnnnnCCGACnnnnnnnnnnnnnnnnnnnnnnnnnnnn ↑

↓ Cut with Fok I

nnnnnGGATGnnnnnnnnnnnnnnnnnnnnnnnnnnnn
 nnnnnCCGACnnnnnnnnnnnnnnnnnnnnnnnnnnnn

Fsp I

↓
 nnnnnnnTGCGCAnnnnnnn
 nnnnnnnACGCGTnnnnnnn ↑

↓ Cut with Fsp I

nnnnnnnTGC GCAnnnnnnn
 nnnnnnnACG CGTnnnnnnn

FIG. 2

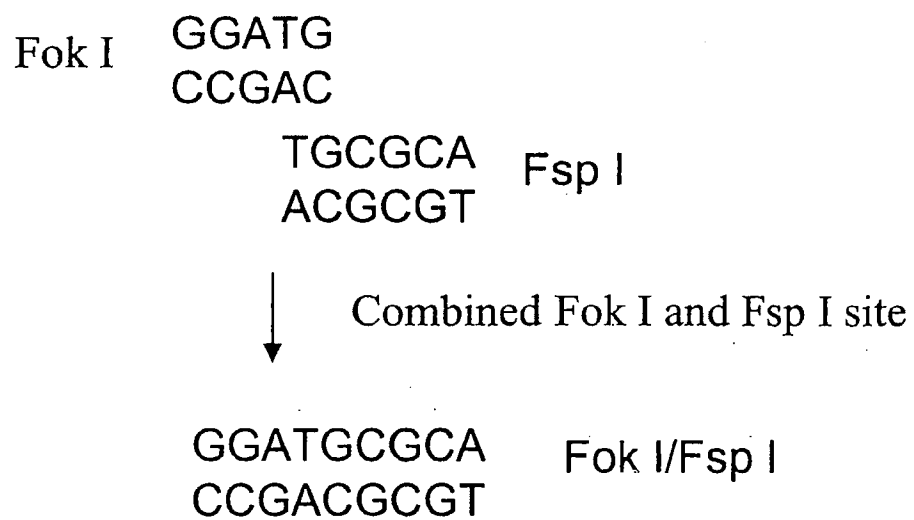


FIG. 3

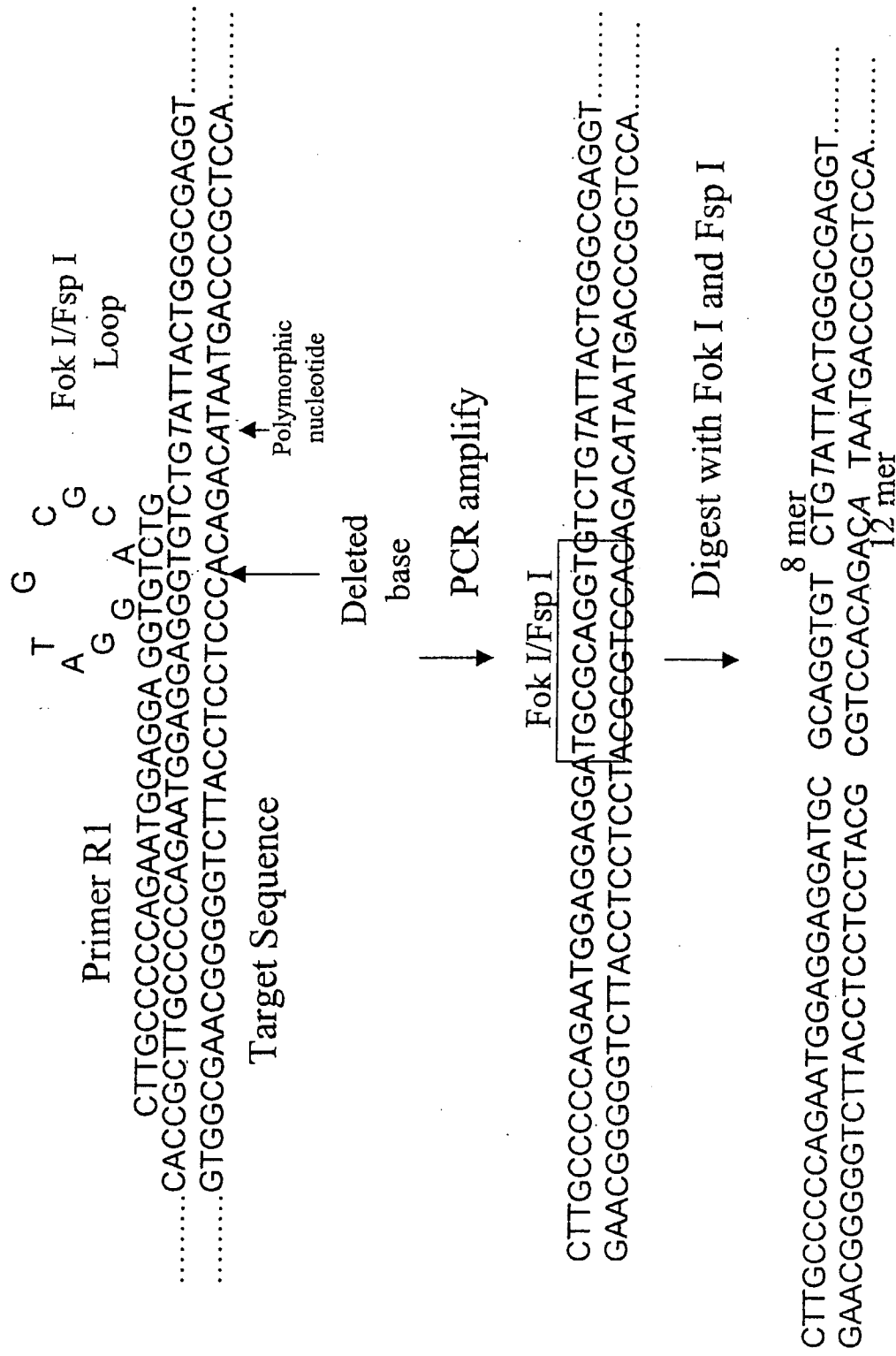


FIG. 4

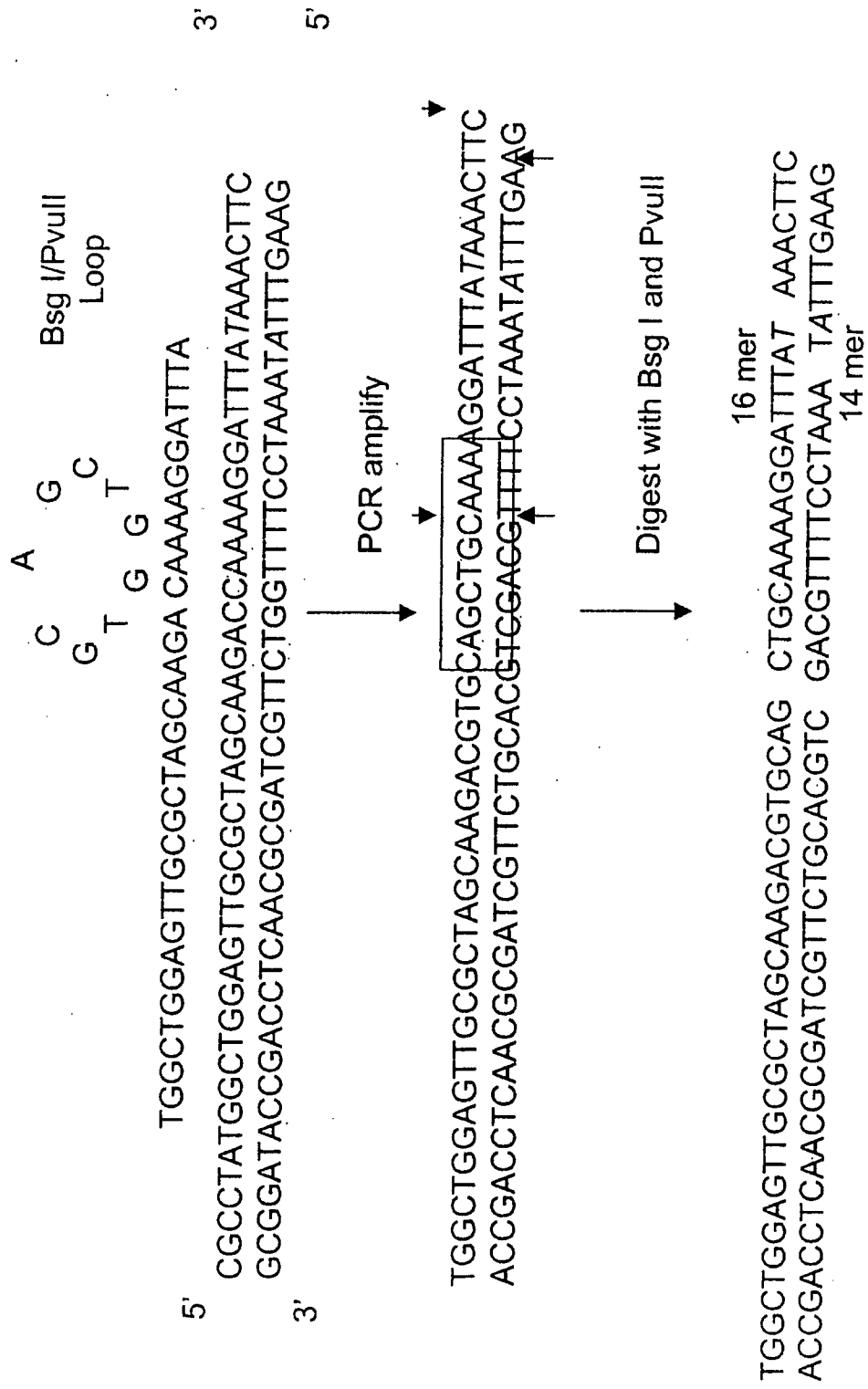


FIG. 5

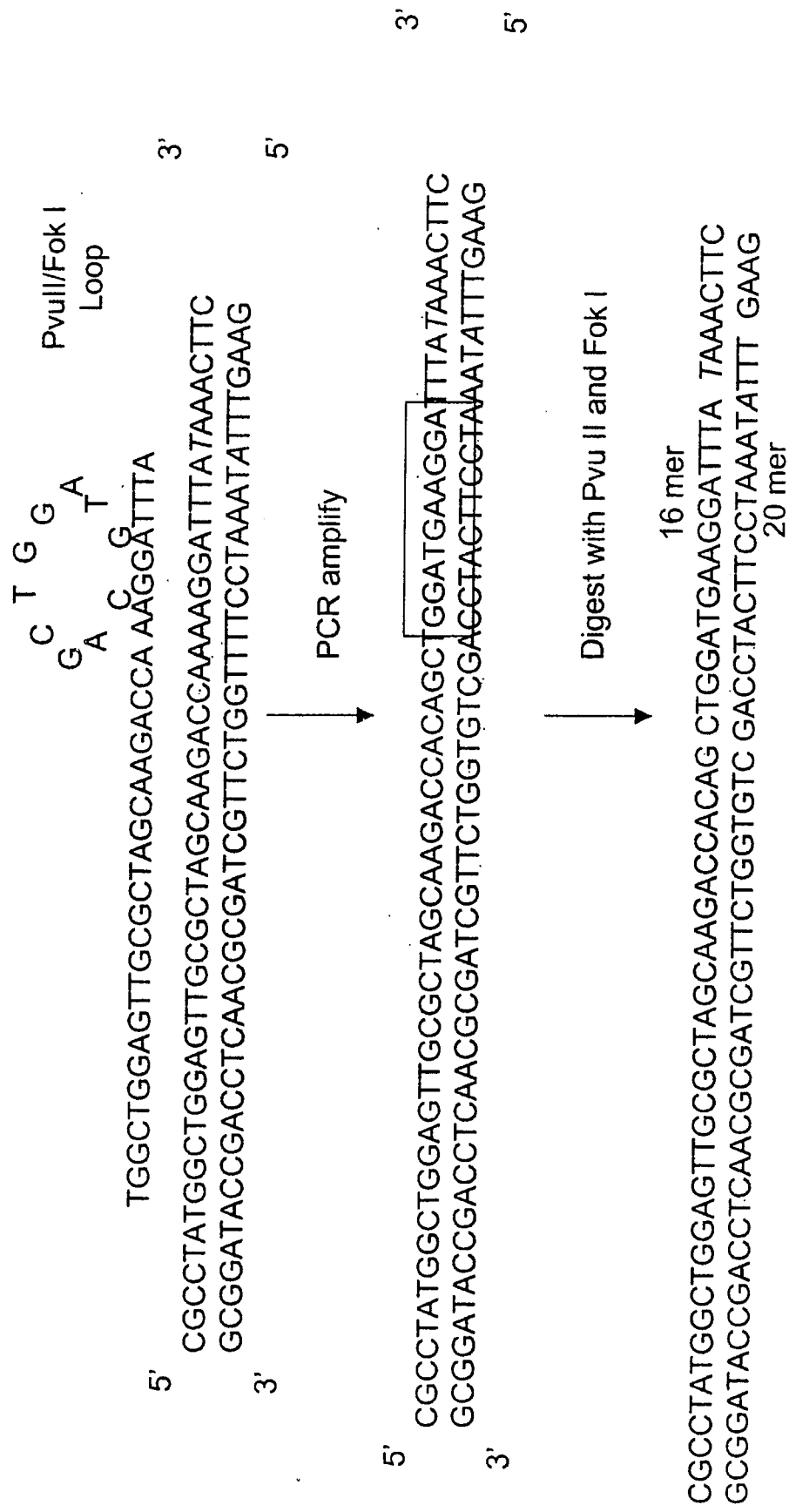


FIG. 6

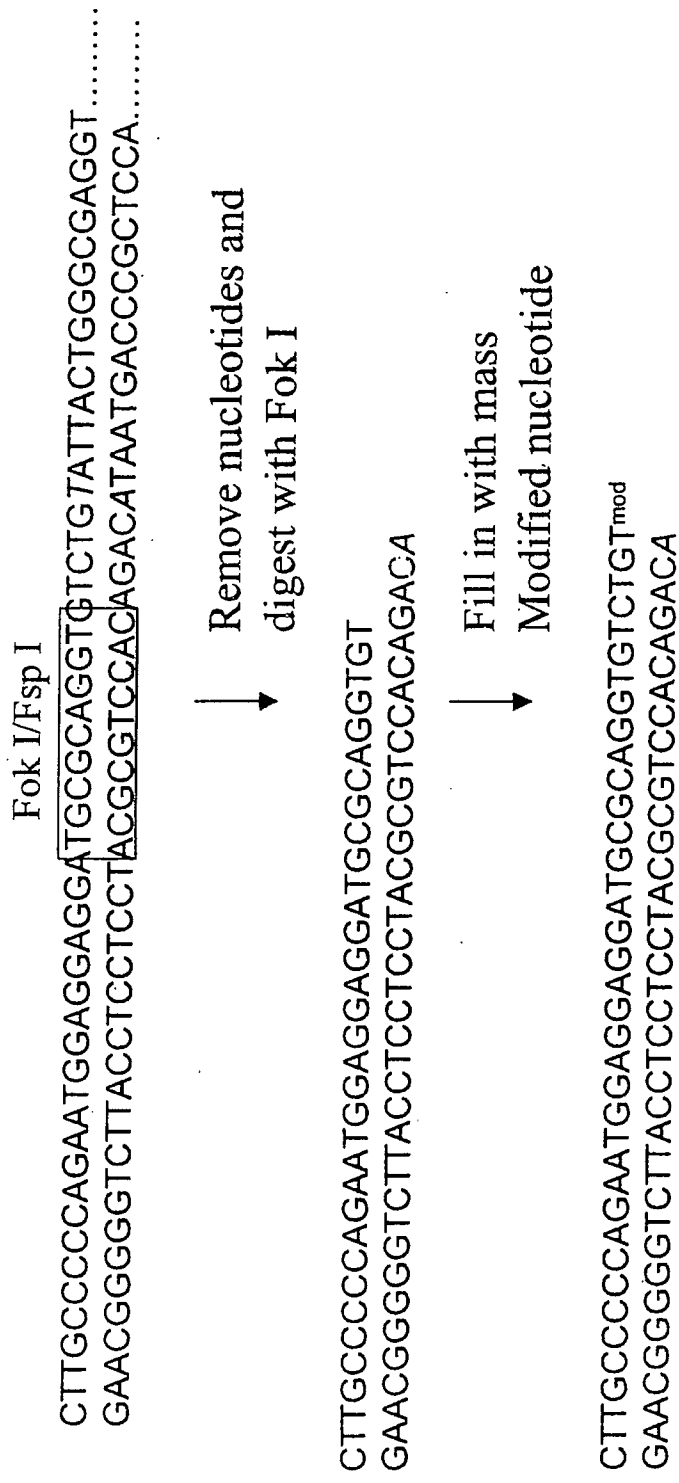


FIG 7

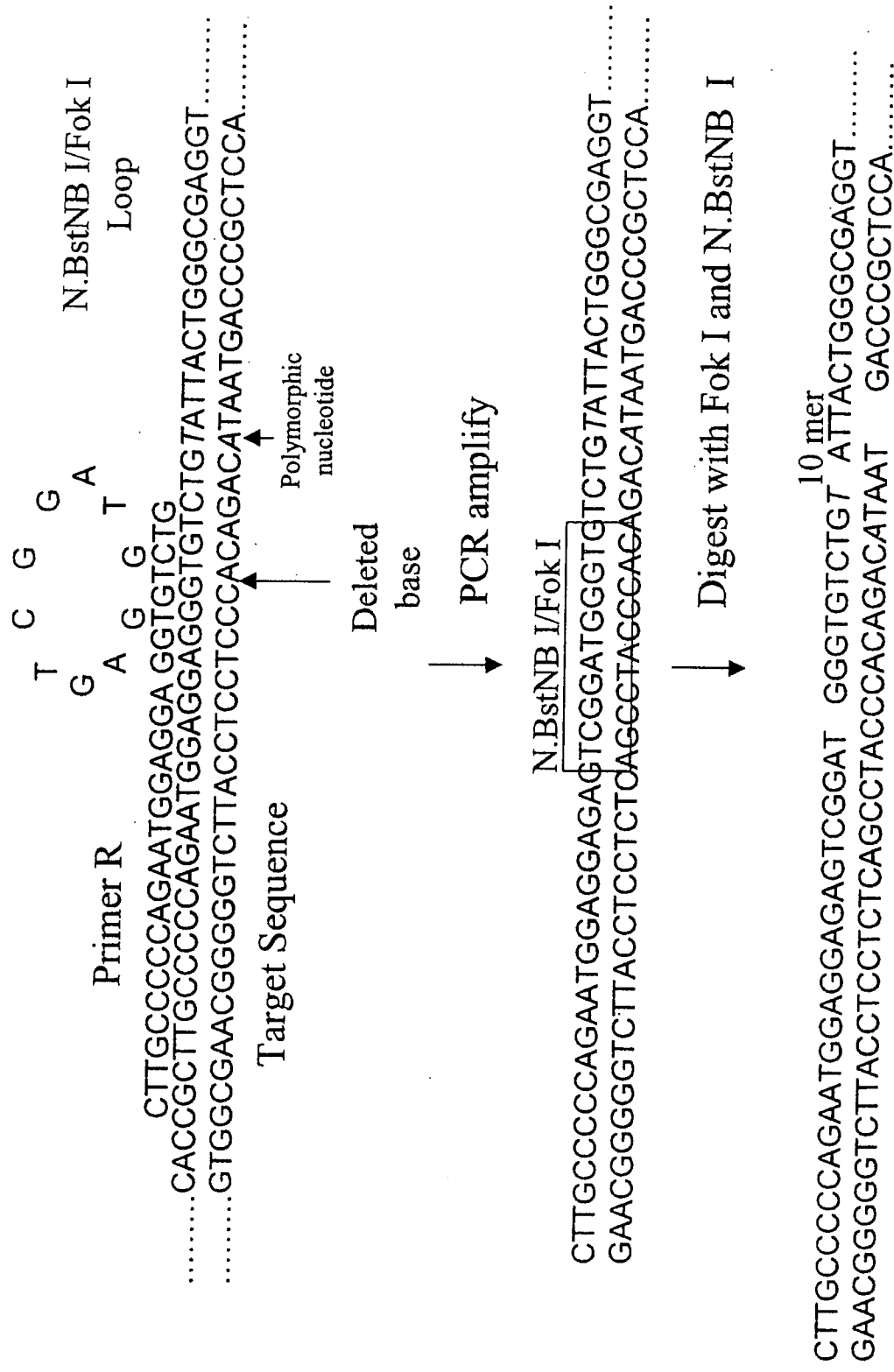


FIG. 9

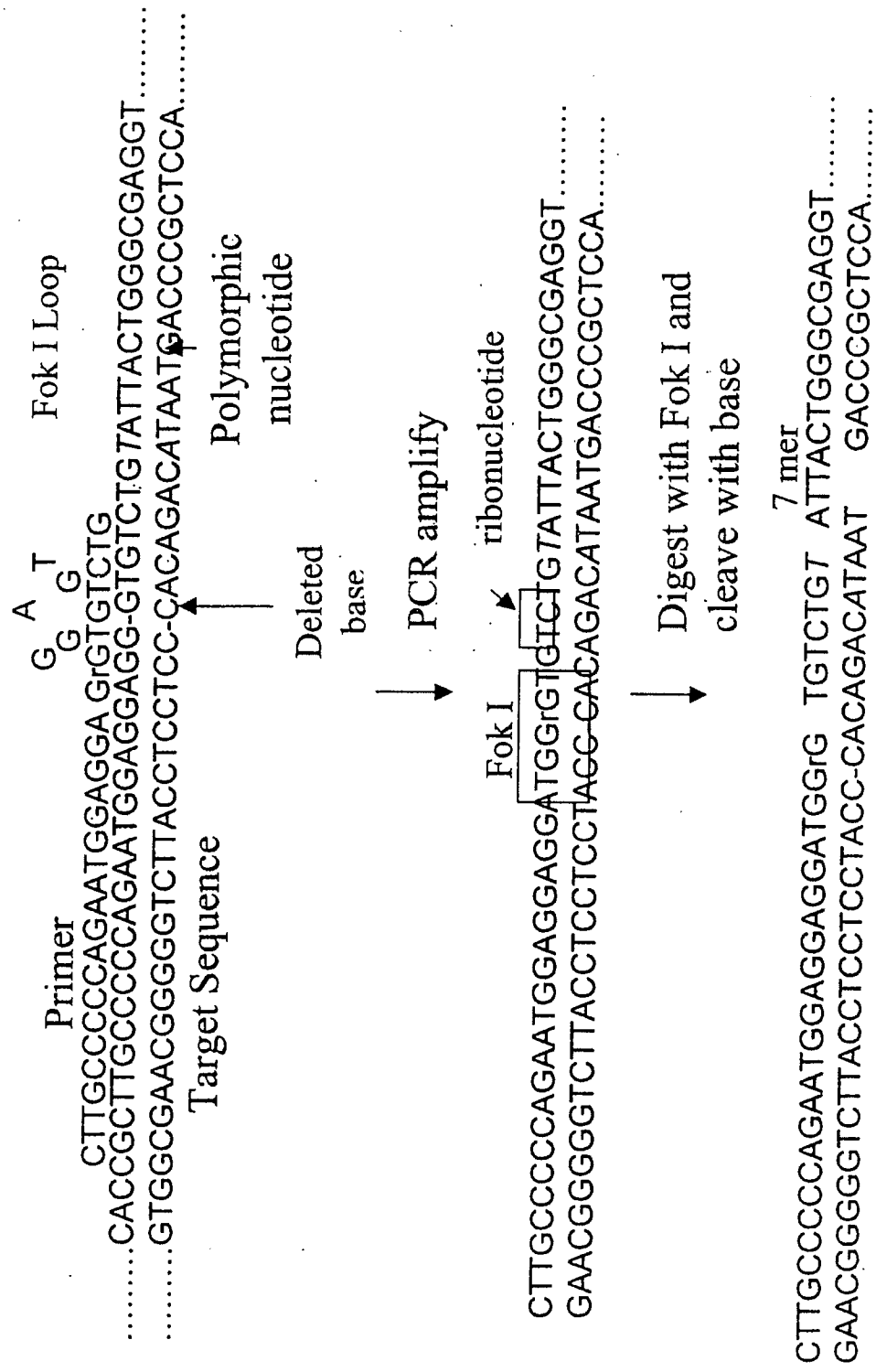


FIG. 10

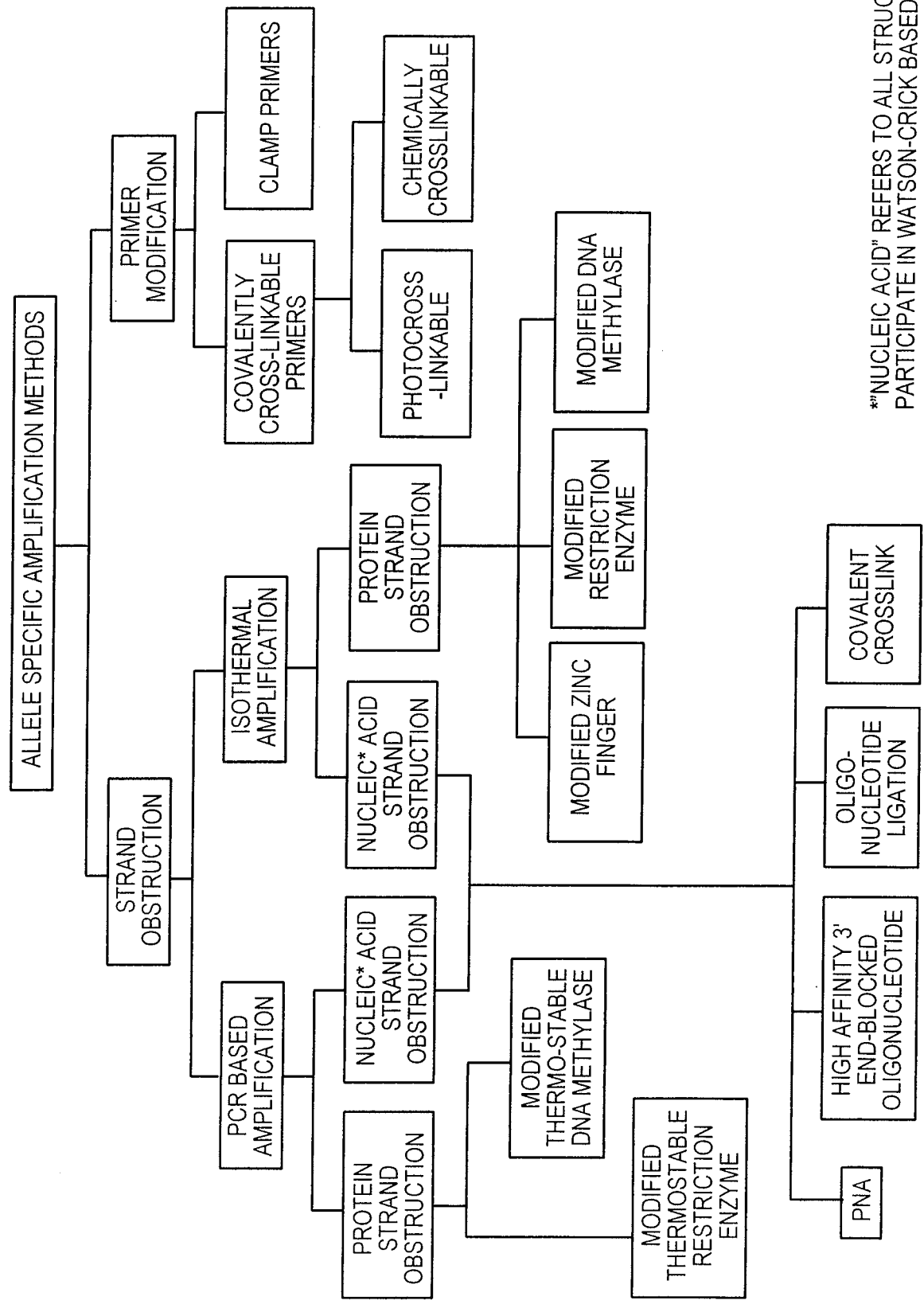
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graph TD
    A[ALLELE SEPARATION METHODS] --> B[SINGLE STRAND SEPARATION METHODS]
    A --> C[DOUBLE STRAND SEPARATION METHODS]
    
    B --> D[PROTEIN AFFINITY CAPTURE]
    B --> E[NUCLEIC ACID AFFINITY CAPTURE]
    
    D --> F[MODIFIED RESTRICTION ENZYME CAPTURE]
    D --> G[PNA BASED CAPTURE]
    D --> H[MODIFIED NUCLEOTIDE CAPTURE]
    D --> I[LIGATION BASED CAPTURE]
    D --> J[COVALENT CROSSLINK CAPTURE]
    D --> K[MODIFIED ZINC FINGER CAPTURE]
    D --> L[MODIFIED RESTRICTION ENZYME CAPTURE]
    
    E --> M[NUCLEIC ACID AFFINITY CAPTURE]
    E --> N[MODIFIED DNA METHYLASE CAPTURE]
    
    C --> O[RESTRICTION SITE POLYMORPHISM BASED METHODS]
    C --> P[HOOGSTEEN BASEPAIRING]
    C --> Q[REVERSE HOOGSTEEN BASEPAIRING]
    
    O --> R[SIZE FRACTIONATION]
    O --> S[END CAPTURE]
    
    R --> T[MODIFIED RESTRICTION ENZYME CAPTURE]
    R --> U[MODIFIED NUCLEOTIDE CAPTURE]
    R --> V[LIGATION BASED CAPTURE]
    R --> W[COVALENT CROSSLINK CAPTURE]
    R --> X[MODIFIED ZINC FINGER CAPTURE]
    R --> Y[MODIFIED RESTRICTION ENZYME CAPTURE]
    R --> Z[MODIFIED DNA METHYLASE CAPTURE]
    
    S --> AA[POLYAMIDE CAPTURE]
    S --> AB[TRIPLEX CAPTURE]
    
    AA --> AC[HOOGSTEEN BASEPAIRING]
    AA --> AD[REVERSE HOOGSTEEN BASEPAIRING]

```

Fig. 11

METHODS FOR HAPLOTYPE ALLELE SPECIFIC AMPLIFICATION



*"NUCLEIC ACID" REFERS TO ALL STRUCTURES THAT PARTICIPATE IN WATSON-CRICK BASED PAIRING

FIG. 12

METHODS FOR HAPLOTYPING BASED ON ALLELE SPECIFIC RESTRICTION

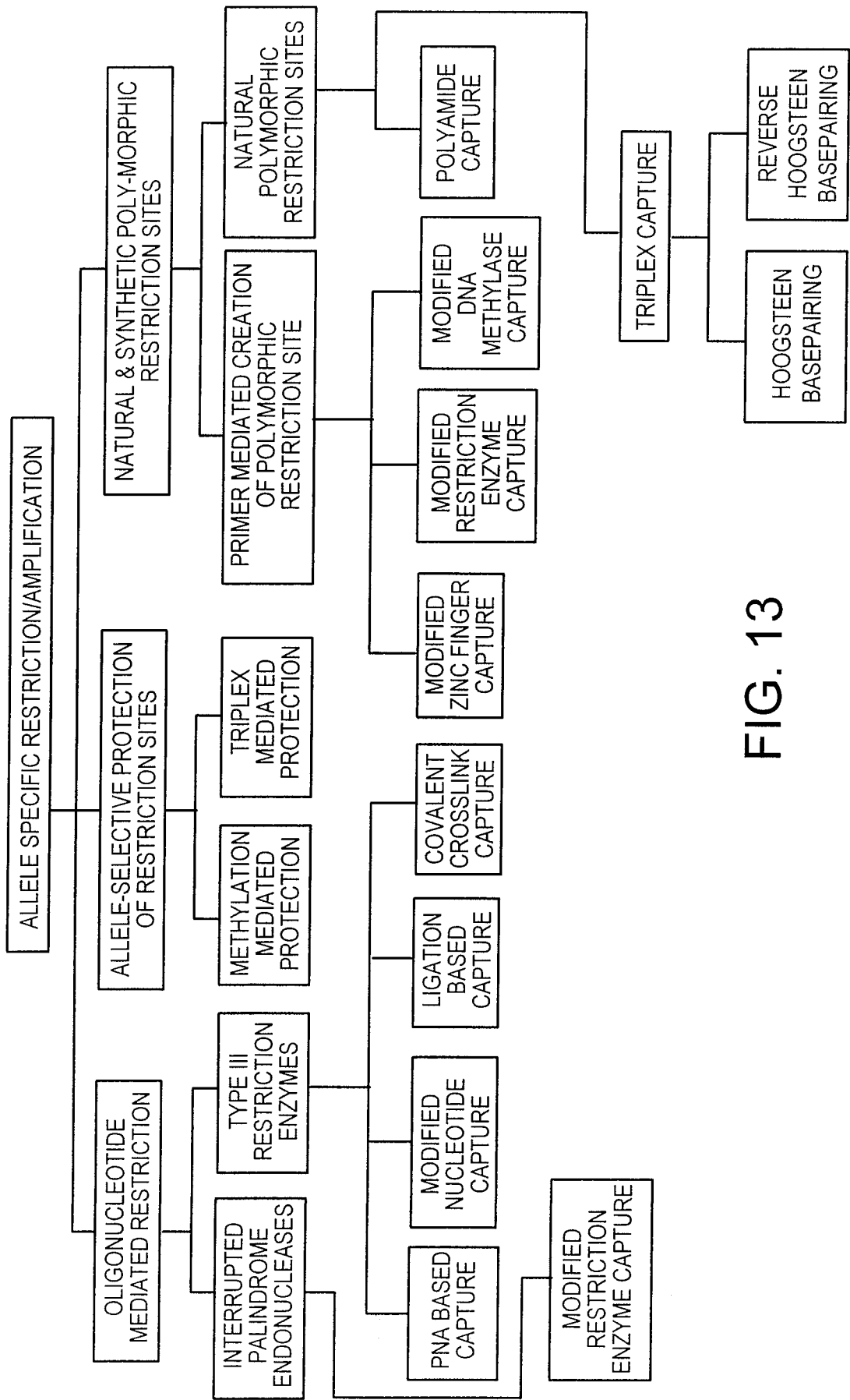


FIG. 13

Hair PCR Primers

ATCTGGANNNNNNNNNNNTCC
____ AGGCTA _____

ALLELE 1
T PRIMER ↓ PCR Amplify

ATCTGGANNNNNNNNNTCCAGAT _____
TAGACCTNNNNNNNNNAGGCTA _____

ATCTGGANNNNNNNNNNNTCC
____ AGGCTA _____

ALLELE 2
T PRIMER ↓ PCR Amplify

ATCTGGANNNNNNNNNTCCGGAT _____
TAGACCTNNNNNNNNNAGGCTA _____

FIG. 14

Hair PCR Primers

ATCCGGANNNNNNNNNNTCC
_____ AGGTCTA _____

ALLELE 1
C PRIMER ↓ PCR Amplify

ATCCGGANNNNNNNNTCCAGAT _____
TAGGCCTNNNNNNNNNAGGTCTA _____

ATCCGGANNNNNNNNNNTCC
_____ AGGCCTA _____

ALLELE 2
C PRIMER ↓ PCR Amplify

ATCCGGANNNNNNNNTCCGGAT _____
TAGGCCTNNNNNNNNNAGGCCTA _____

FIG. 15

Minus strand resulting from PCR of allele 1

Minus Strand

↓

Hairpin loop forms inhibiting hybridization of PCR primer and amplification of allele 1

ALLELE 1
T PRIMER

Minus Strand

FIG. 16

The diagram illustrates the principle of Allele-Specific PCR (AS-PCR) for genotyping. It compares two scenarios:

- Allele 1:** The target sequence is TAGGCCTNNNNNNNAGGTCTA. The primer used is NNNNNNNNTCCGAT. There is a mismatch at the 3' end (the primer has a G where the template has a T). This mismatch prevents the primer from hybridizing correctly, inhibiting PCR amplification.
- Allele 2:** The target sequence is TAGACCTNNNNNNNAGGCCTA. The primer used is NNNNNNNNTCCGGAT. This primer perfectly matches the 3' end of the target sequence, allowing for successful hybridization and subsequent PCR amplification.

FIG. 17

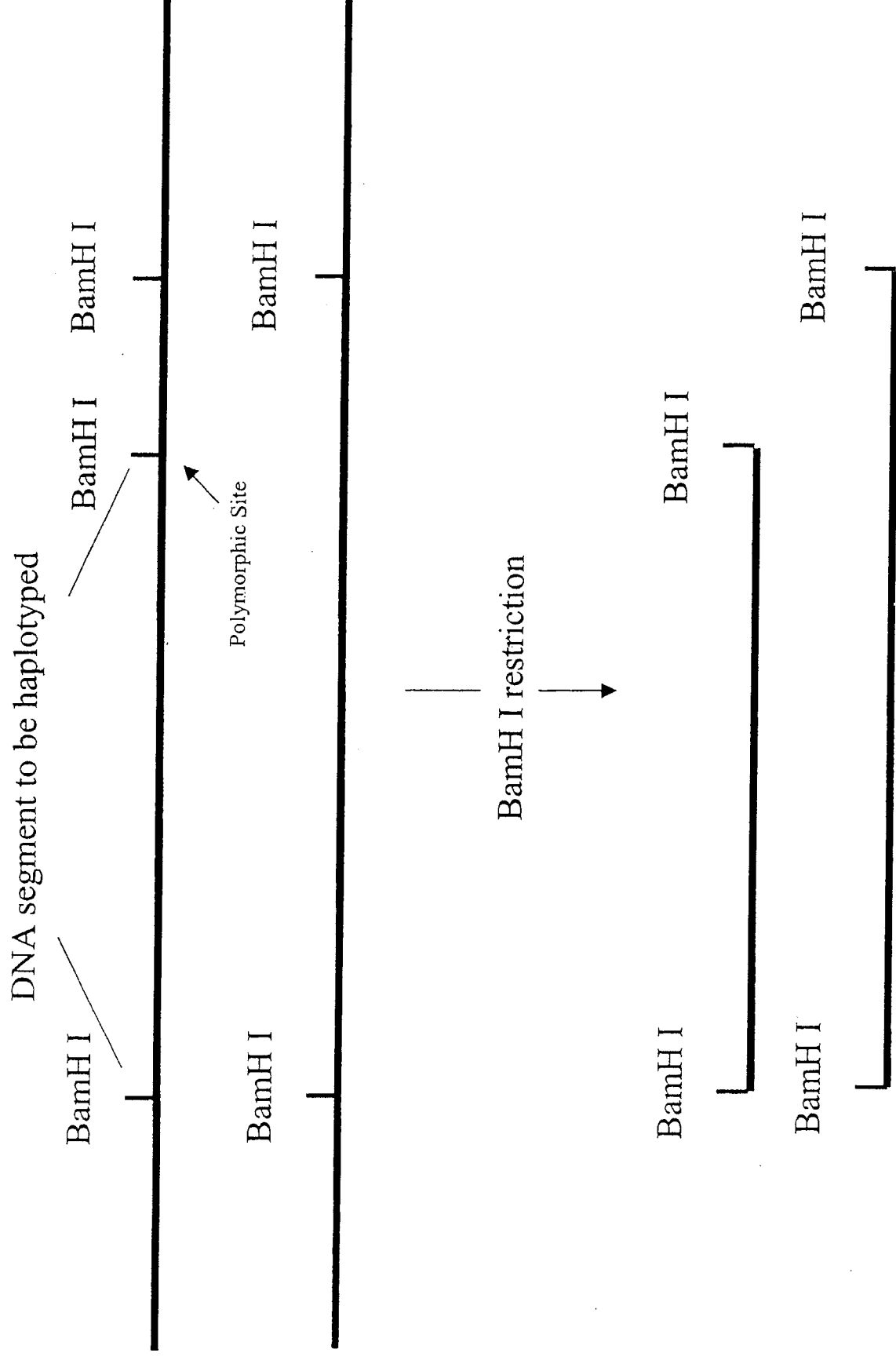


FIG 18

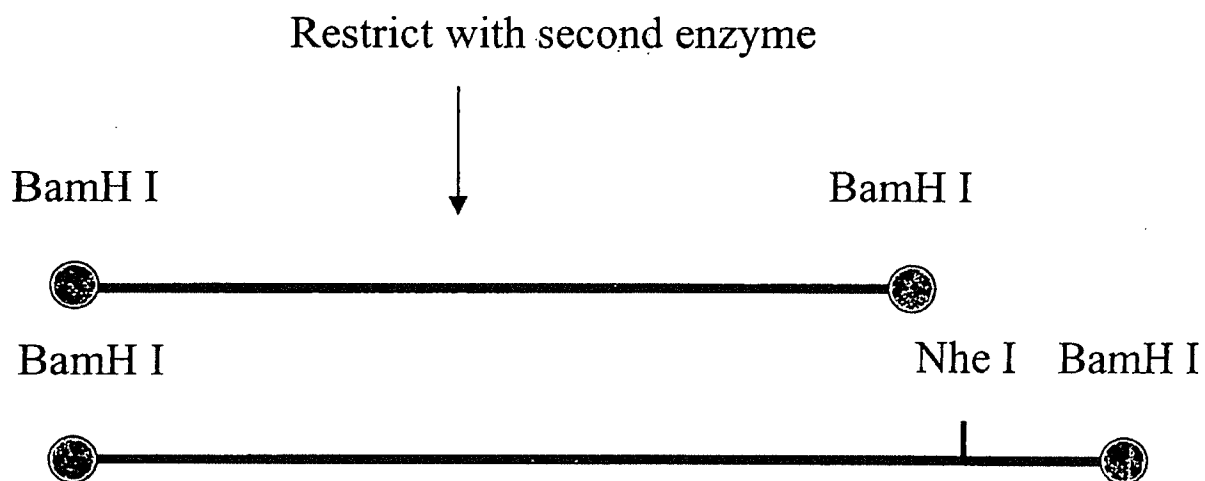
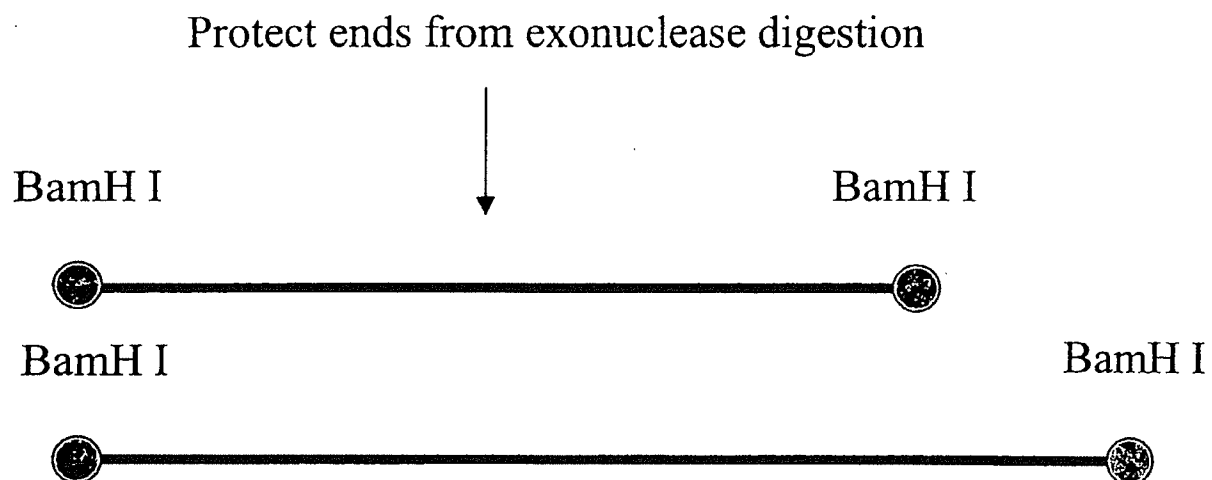


FIG. 19

Digest with exonuclease

Add single strand nuclease to remove/degrade remaining single strand

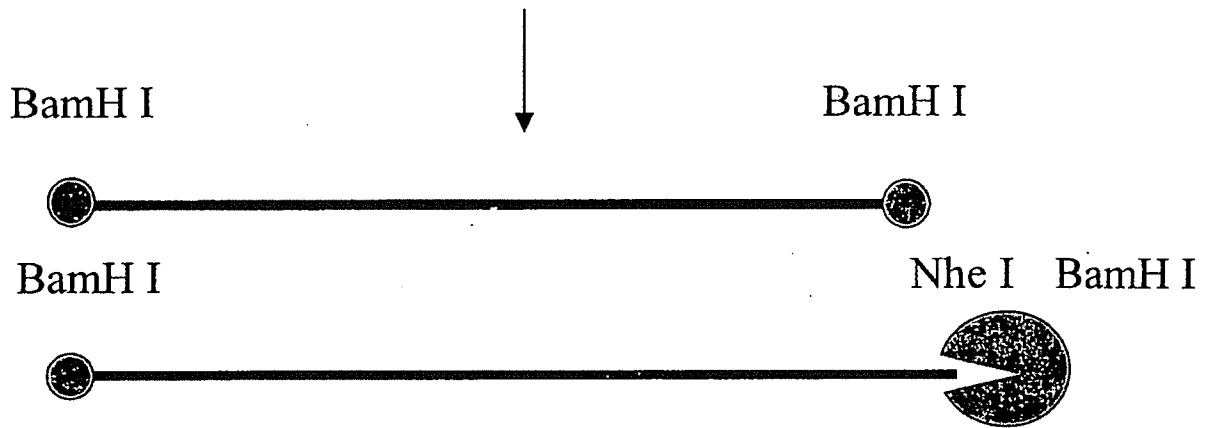


FIG. 20

Dihydropyrimidine dehydrogenase (DPD) polymorphisms used in haplotyping assay.

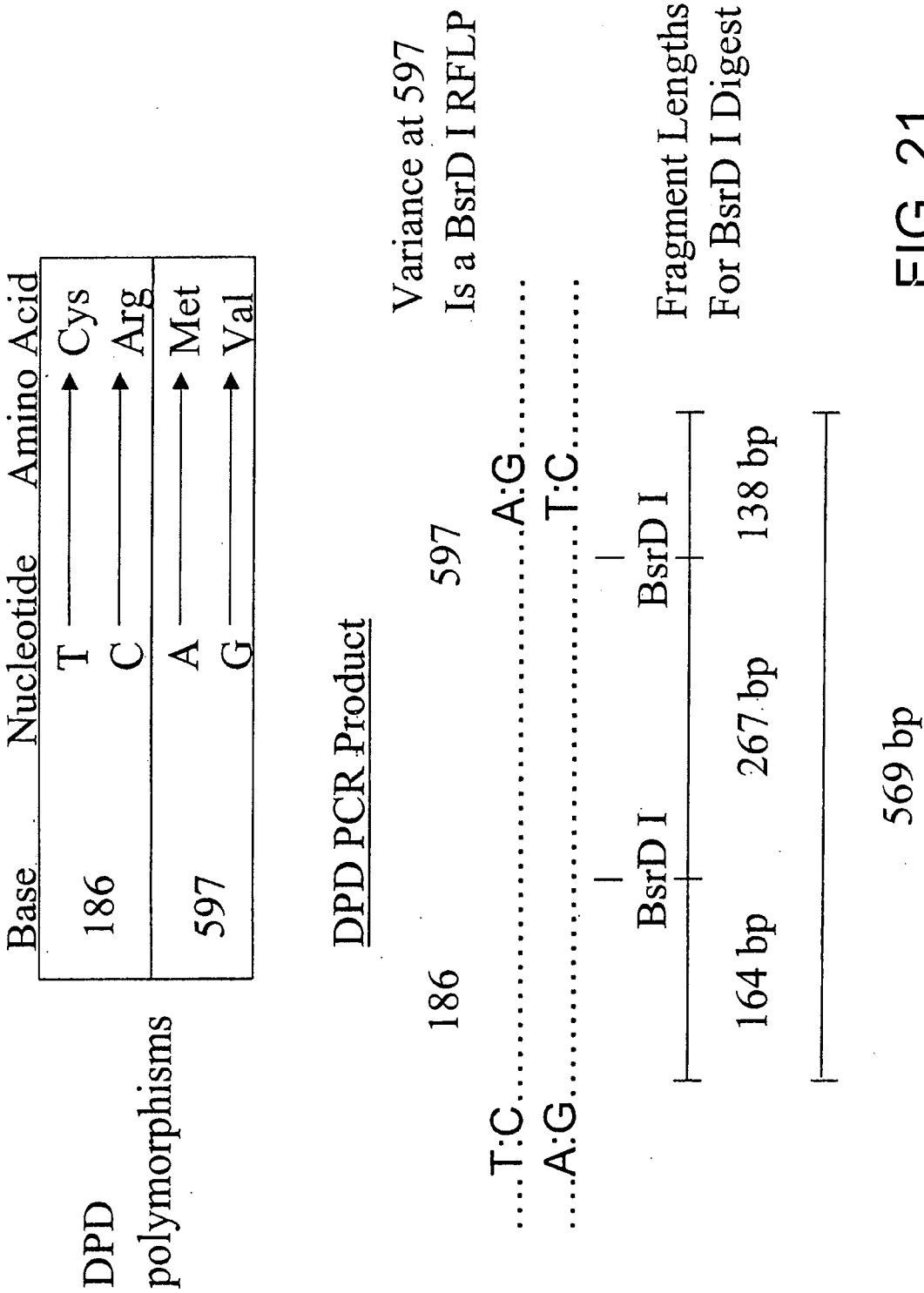


FIG. 21

Allele Specific Primers for DPD

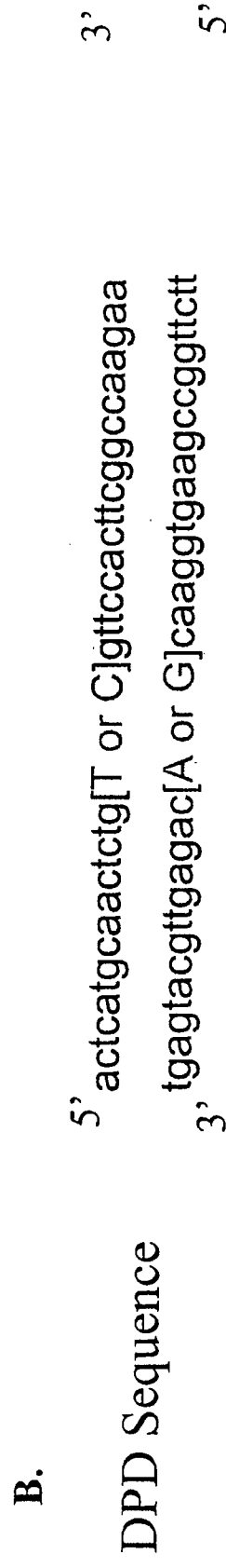
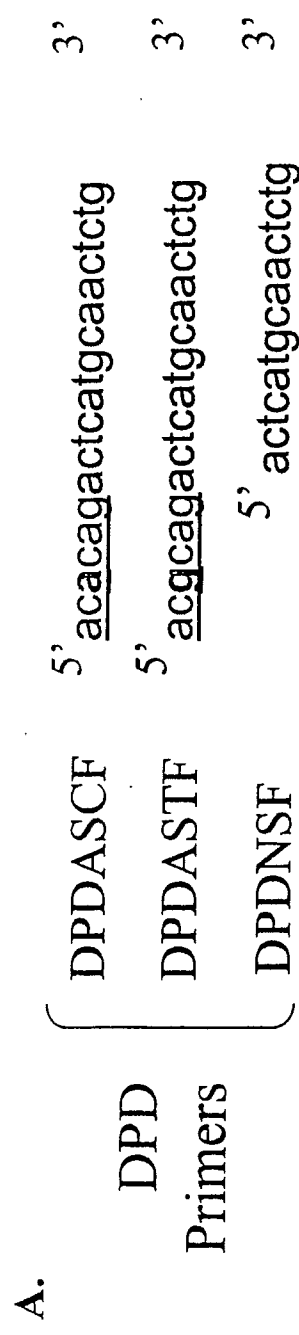


FIG. 22

PCR Amplification Using DPDNSF Primer

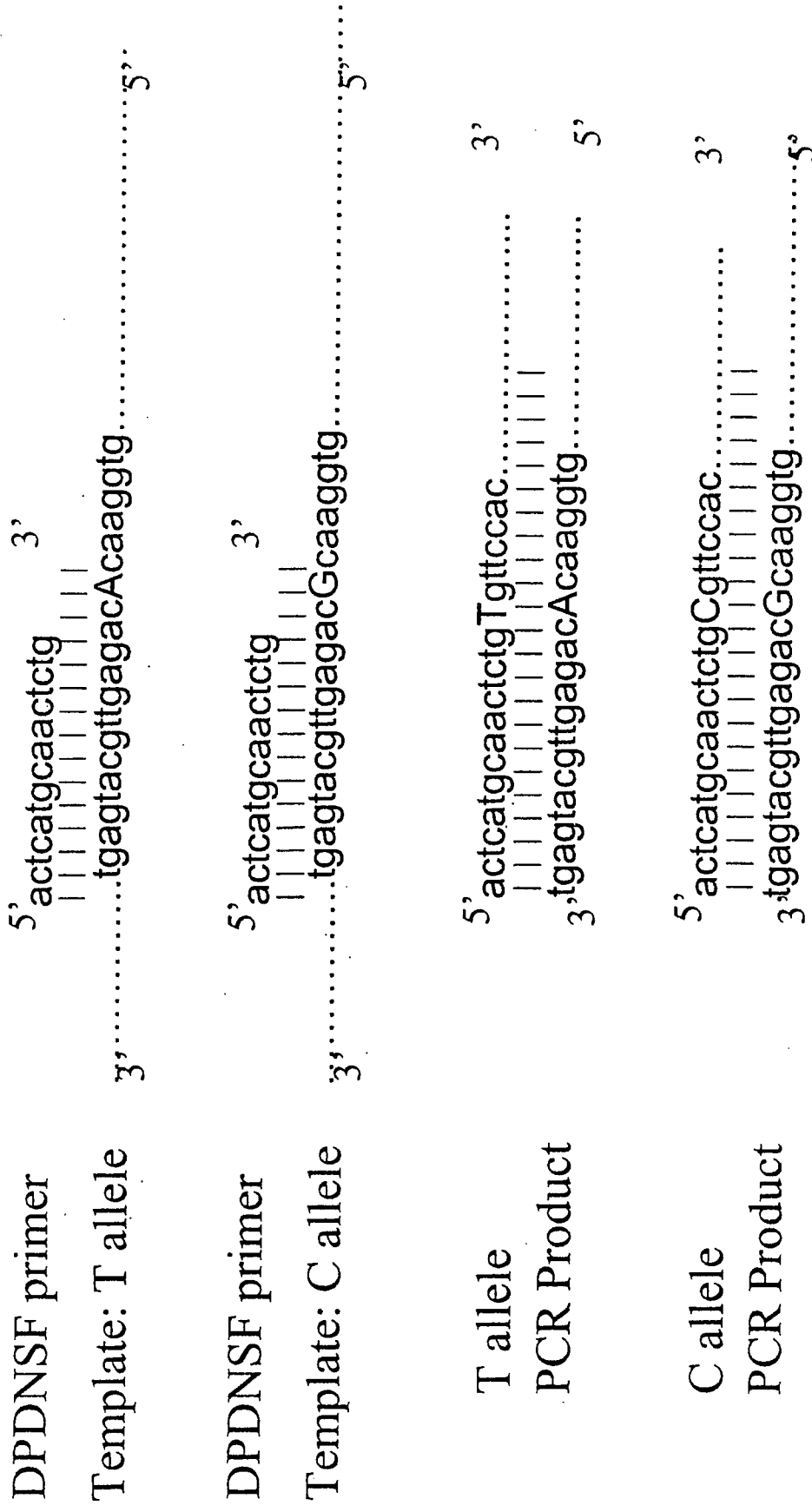


FIG. 23

PCR Amplification Using DPDASTF Primer

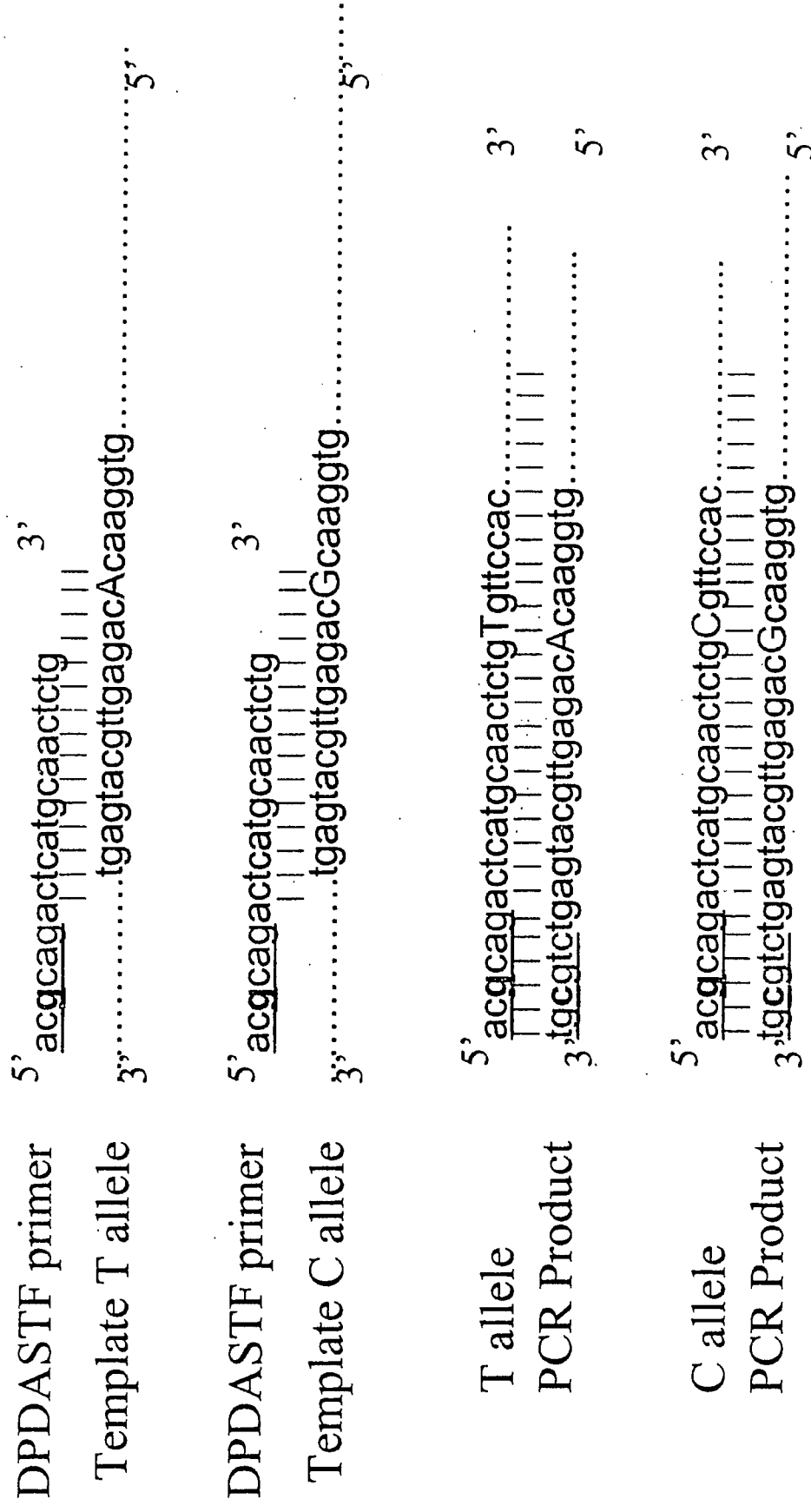


FIG. 24

PCR Amplification Using DPDASCF Primer

DPDASCF primer

5' acacagactcatgcaactctg 3'

Template T allele

3'.....tgagtacgttgagacAcaagggtg.....5'

DPDASCF primer

5' acacagactcatgcaactctg 3'

Template C allele

3'.....tgagtacgttgagacGcaagggtg.....5'

T allele

5' acacagactcatgcaactctgTgttccac..... 3'

PCR Product

3'-tgtgtctgagtacgttgagacAcaagggtg..... 5'

C allele

5' acacagactcatgcaactctgCgttccac..... 3'

PCR Product

3'-tgtgtctgagtacgttgagacGcaagggtg..... 5'

FIG. 25

Hairpin Structures for PCR Products Generated Using DPDNSF Primer

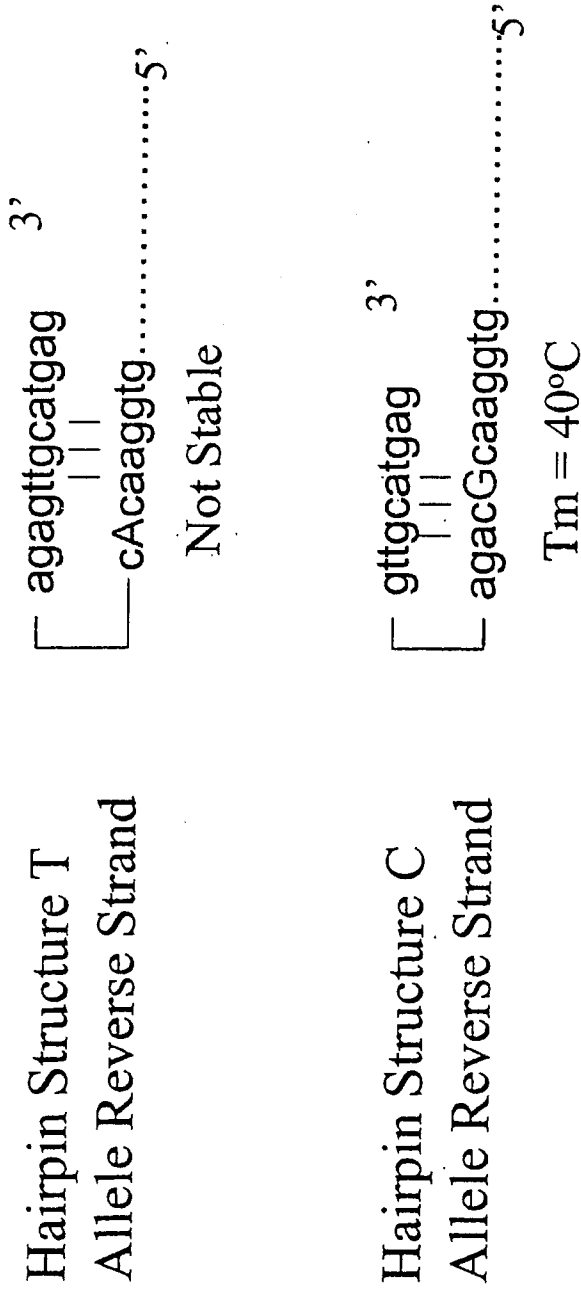


FIG. 26

Hairpin Structures for PCR Products Generated Using WPDASCF Primer

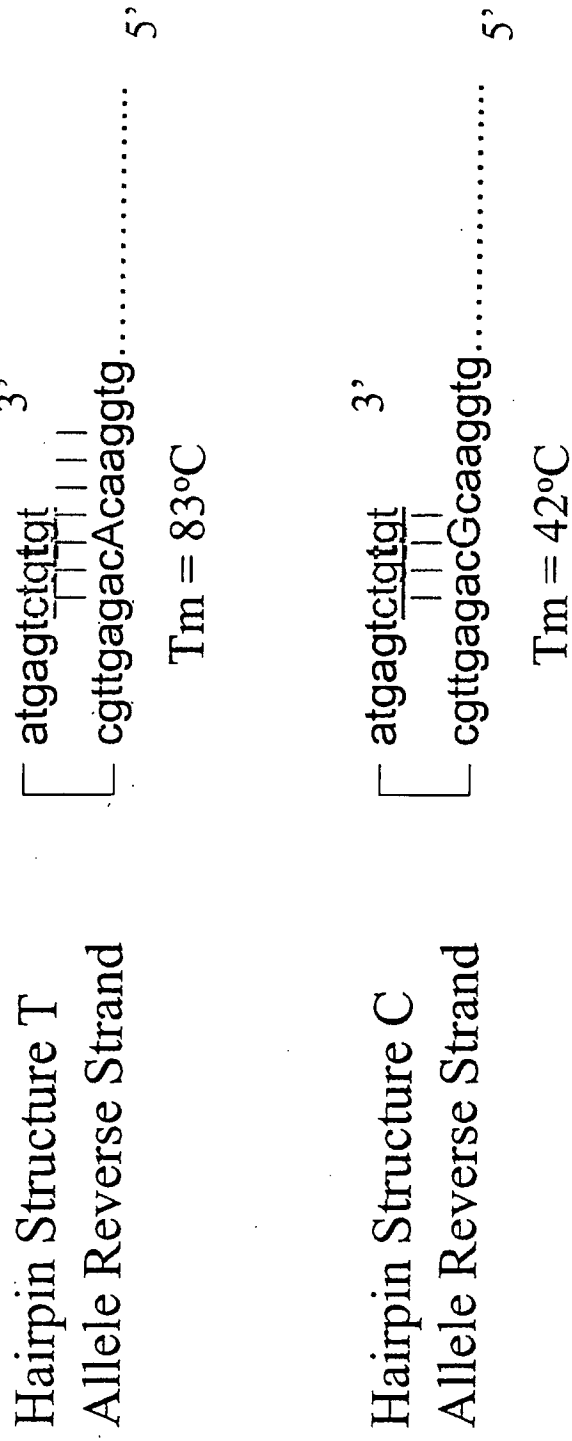


FIG. 27

Hairpin Structures for PCR Products Generated Using DPDASTF Primer

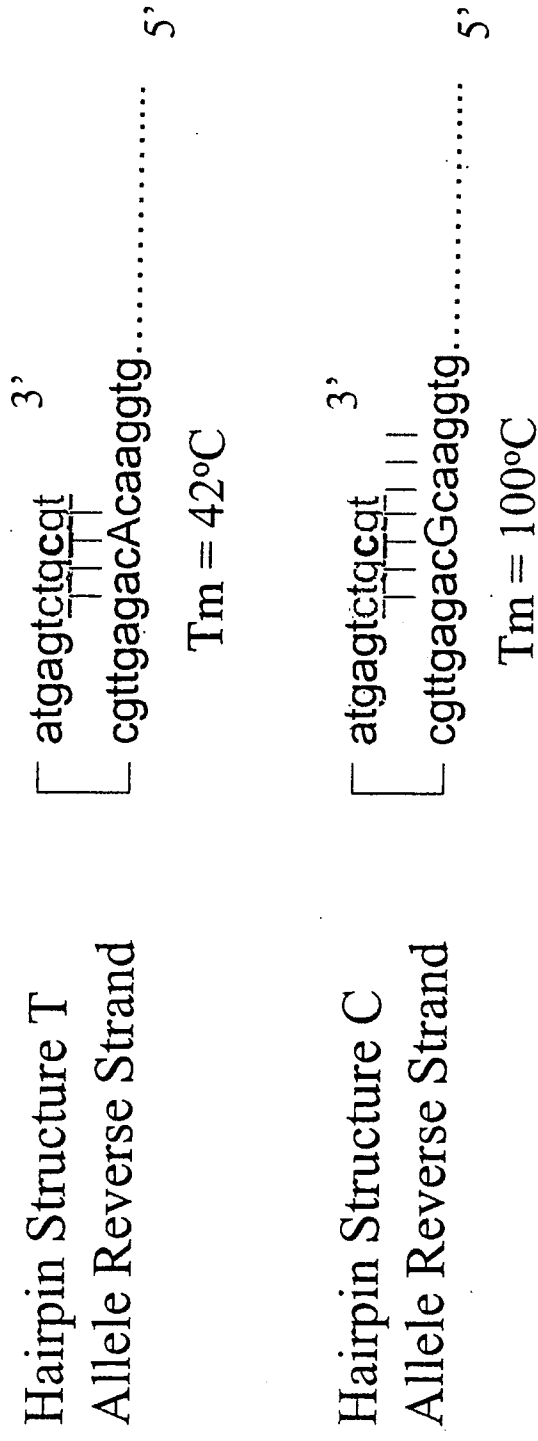


FIG. 28

Non-Allele Specific Amplification Using DPDNSF Primer

ALLELE C

DPDNSF primer

5' actcatgcaactctg 3' T_m = 41°C

[gttgcatgag 3'
 |||
 [agacGcaagtg.....5' T_m = 40°C

↓
 Primer
 Hybridization
 and Amplification

5' actcatgcaactctg 3'
 |||
 3' tgagtacgttgagacGcaagtg... 5'

ALLELE T

DPDNSF primer

5' actcatgcaactctg 3' T_m = 41°C

[agagttgcatgag 3'
 |||
 [cAcaaggtg.....5' Not Stable

↓
 Primer
 Hybridization
 and Amplification

5' actcatgcaactctg 3'
 |||
 3' tgagtacgttgagacAcaagtg... 5'

FIG. 29

Allele Specific Amplification Using DPDASCF Primer

ALLELE C

DPDASCF primer T_m = 60°C
 5' acacagactcatgcaactctg 3'

atgagtcctgtgt
 ||||
 [cgttgagacGcaagggtg..... 5'

↓
 Primer
 Hybridization
 and Amplification

5' acacagactcatgcaactctg 3'
 ||||
 3' tgtgctgagtcgttgagacGcaagggtg... 5'

ALLELE T

DPDASCF primer T_m = 60°C
 5' acacagactcatgcaactctg 3'

atgagtcctgtgt
 ||||
 [cgttgagacAcaagggtg..... 5'

↓
 Hairpin inhibits
 Primer Hybridization
 and Amplification

5' acacagactcatgcaactctg 3'
 atgagtcctgtgt
 ||||
 5' [cgttgagacAcaagggtg..... 5'

FIG 30

Allele Specific Amplification Using DPDASTF Primer

ALLELE C

DPDASTF primer T_m = 65°C

5' acgcagactcatgcaactctg

[atgagtctgcgt
|||
cgttgagacGcaagggtg..... T_m = 100°C

↓
Hairpin inhibits
primer hybridization
and Amplification

5' acgcagactcatgcaactctg 3'

[atgagtctgcgt
|||
cgttgagacGcaagggtg..... 5' 3'

ALLELE T

DPDASTF primer T_m = 65°C

acgcagactcatgcaactctg

[atgagtctgcgt
|||
cgttgagacAcaagggtg..... T_m = 42°C 5'

↓
Primer hybridizes
and amplification ensues

5' acgcagactcatgcaactctg 3'
|||
cgttgagacAcaagggtg... 5' 3'

FIG. 31

Allele Specific Amplification of a Heterozygous
Sample with Haplotype T¹⁸⁶, A⁵⁹⁷ and C¹⁸⁶, G⁵⁹⁷

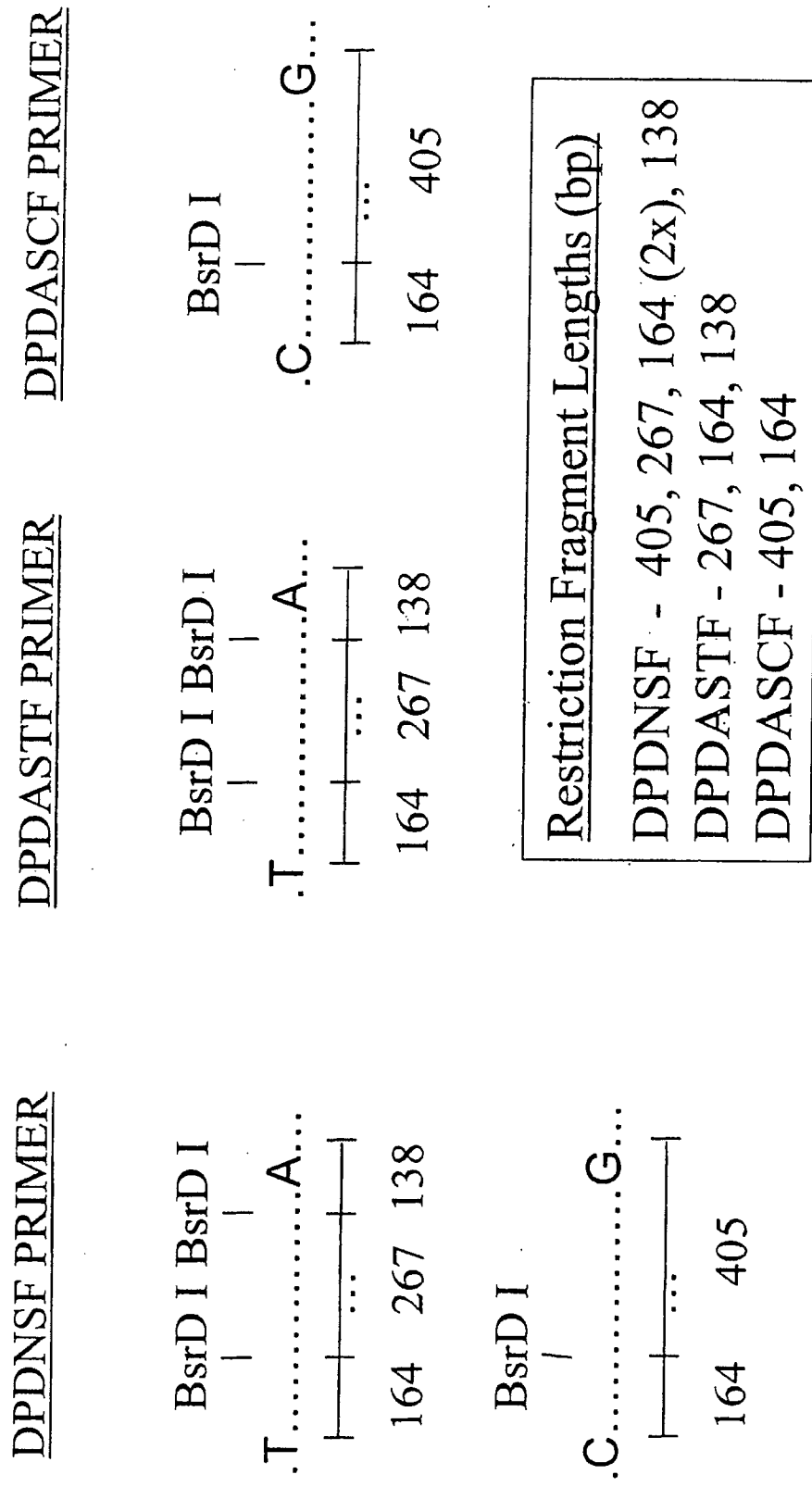


FIG. 32

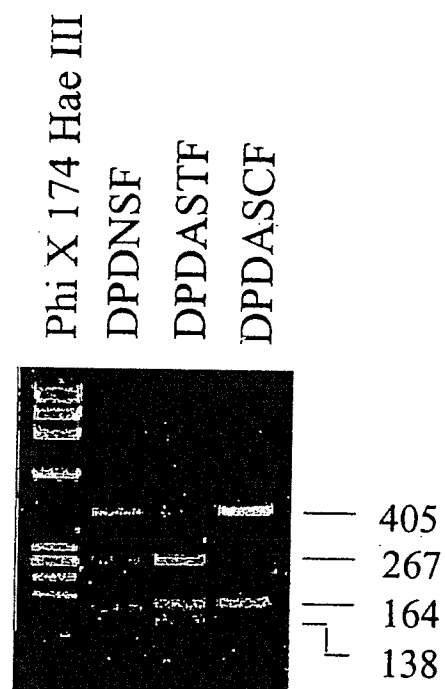


FIG. 33

T Allele Amplicon

↓ ↓
CCCGGCTGGCGCGGACATGGGATGCGCAAGGACGTGTGCGGCCGCCCTGGTGCAGTAC
GGCCGACCCGCGCCTGTACCTACGCGT↓CCTGCACACGCCCGCGGACCAACGTCATG

CGCGGCGAGGTGCAGGCCATGCTCGGCCAGAGACCGAGAGCTGCGGGTGCGCCCTCG
GCGCCGCTCCACGTCCGGTACGAGCCGGTCTCGTGGCTCCTCGACGCCACCGGAGC

CCTCCACCTGCGCAAGCTGCGTAAGCGGCTCCTCCGCGATGCCGATGACCTGCAGAAGC
GGAGGTGACGCGTTCGACGCATTGCCCGAGGAGCGGCTACGGCTACTGGACGCTCTTCG

C Allele Amplicon

↓ ↓
CCCGGCTGGCGCGGACATGGGATGCGCAAGGACGTGCGGCCGCCCTGGTGCAGTAC
GGCCGACCCGCGCCTGTACCTACGCGT↓CCTGCACGCCCGCGGACCAACGTCATG

CGCGGCGAGGTGCAGGCCATGCTCGGCCAGAGACCGAGAGCTGCGGGTGCGCCCTCG
GCGCCGCTCCACGTCCGGTACGAGCCGGTCTCGTGGCTCCTCGACGCCACCGGAGC

CCTCCACCTGCGCAAGCTGCGTAAGCGGCTCCTCCGCGATGCCGATGACCTGCAGAAGC
GGAGGTGACGCGTTCGACGCATTGCCCGAGGAGCGGCTACGGCTACTGGACGCTCTTCG

FIG. 35

[illegible]

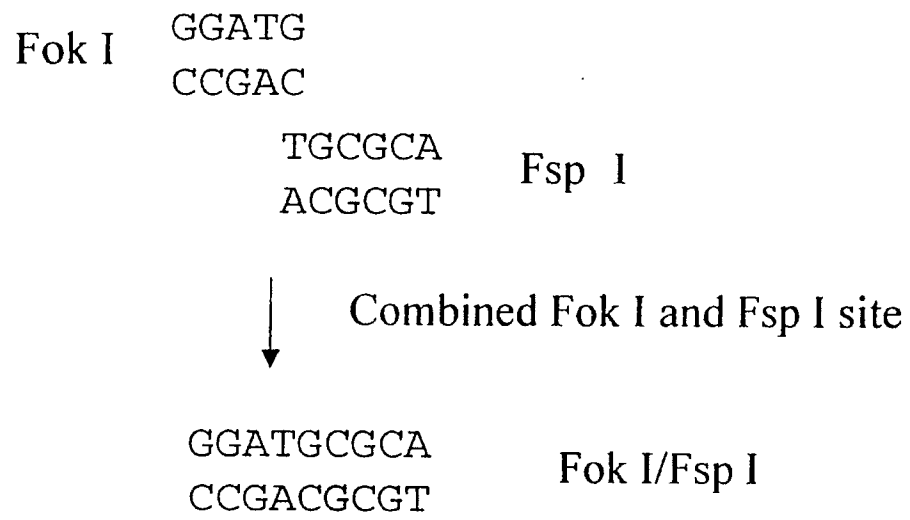


FIG. 3

Restriction Enzyme Genotyping

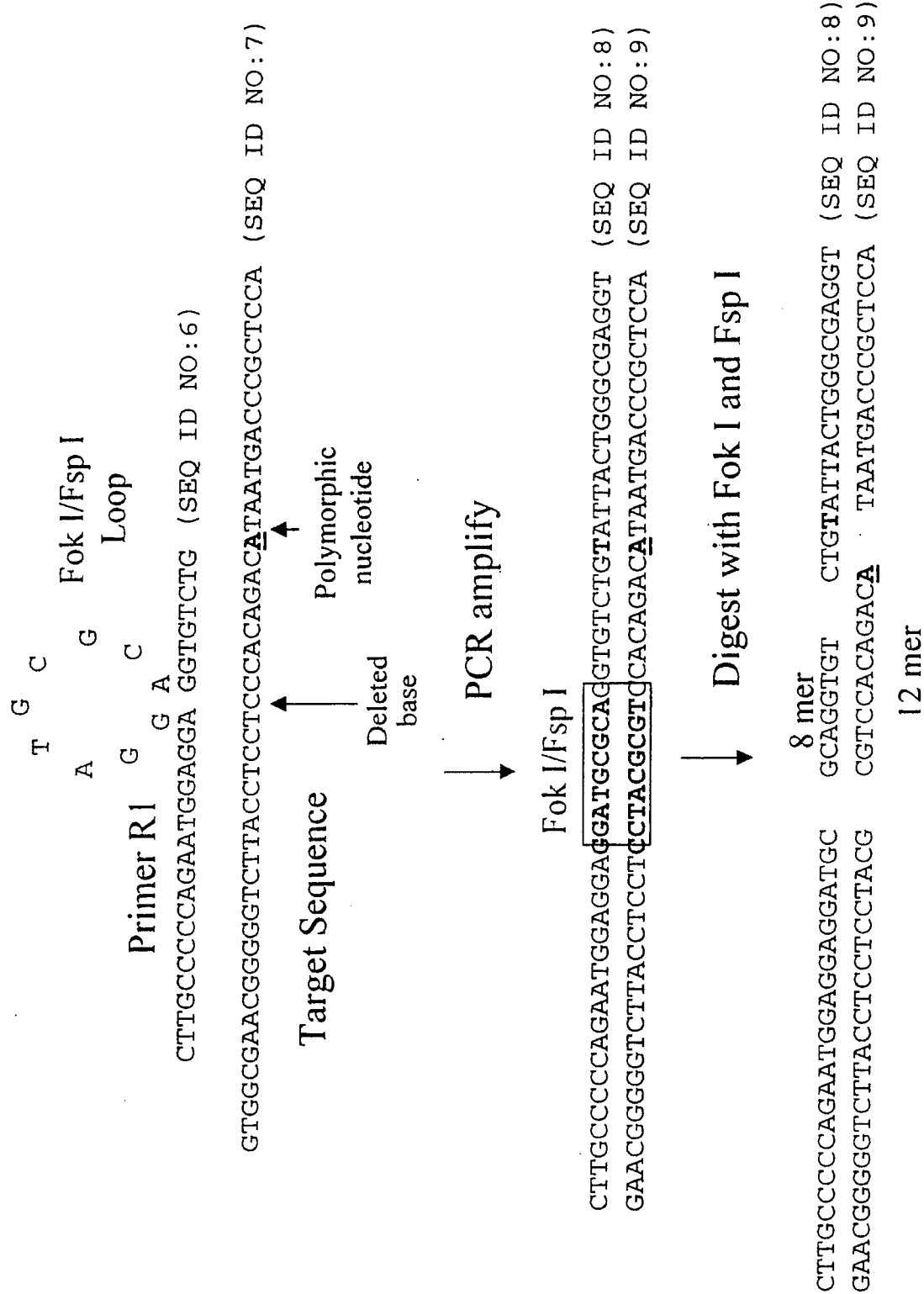


FIG. 4

Introduction of Fok I and Pvu II sites during PCR by loop followed by endonuclease digestion

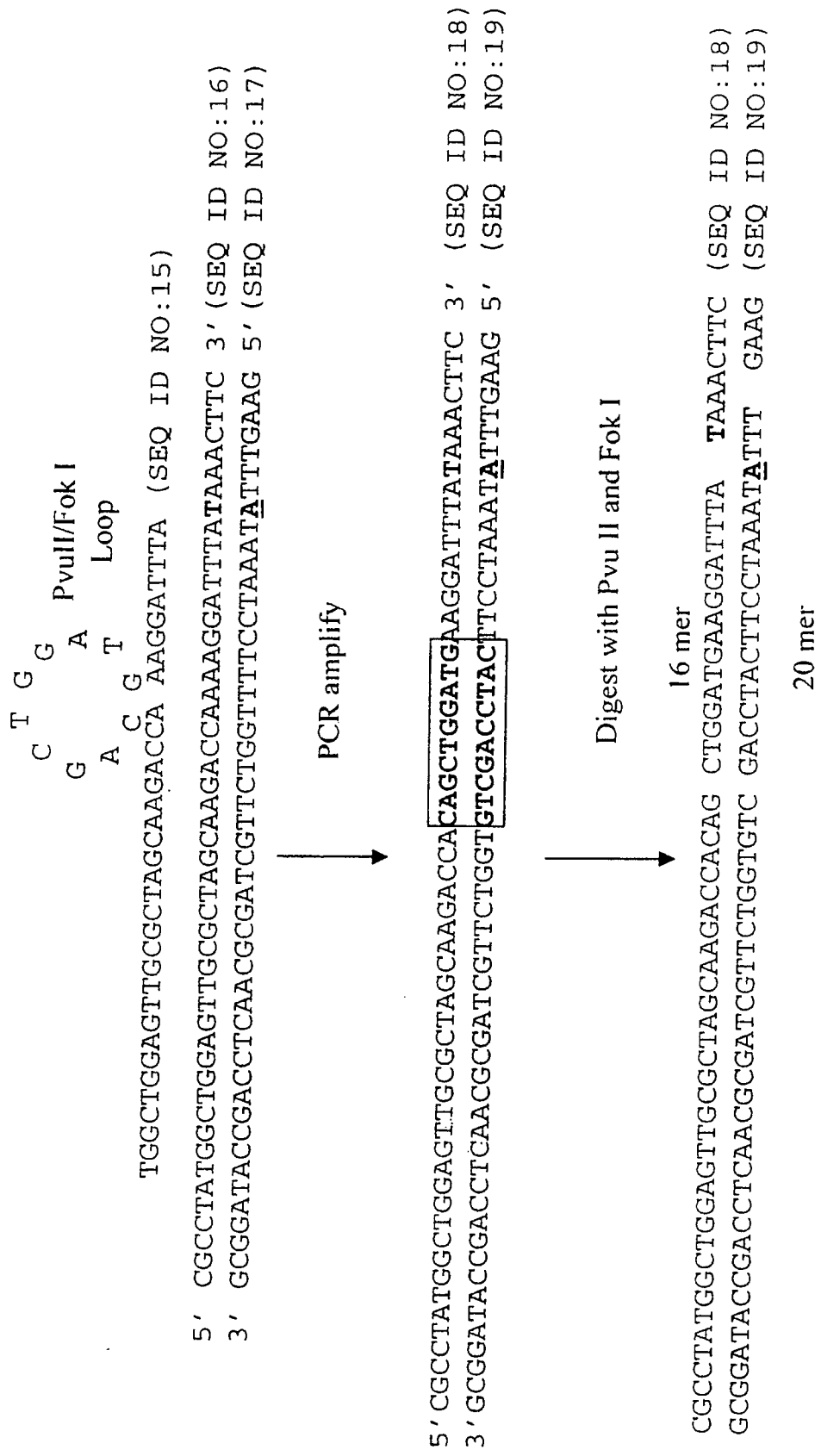


FIG. 6

Fok I/Fsp I

CTTGCCCCCAGAAATGGAGGAGGATGCGCA^{GGTGTCTGTATTACTGGGCGAGGT} (SEQ ID NO:20)
GAACGGGGGTCTTACCTCCTCCTACGCGGTCCACAGACA^{TAAATGACCCGCTCCA} (SEQ ID NO:21)

↓
Remove nucleotides and
digest with Fok I

CTTGCCCCCAGAAATGGAGGAGGATGCGCAGGTGT (SEQ ID NO:22)
GAACGGGGGTCTTACCTCCTCCTACGCGGTCCACAGACA_A (SEQ ID NO:23)

↓
Fill in with mass
Modified nucleotide

CTTGCCCCCAGAAATGGAGGAGGATGCGCAGGTGTCTGT^{mod} (SEQ ID NO:24)
GAACGGGGGTCTTACCTCCTCCTACGCGGTCCACAGACA_A (SEQ ID NO:23)

FIG. 7

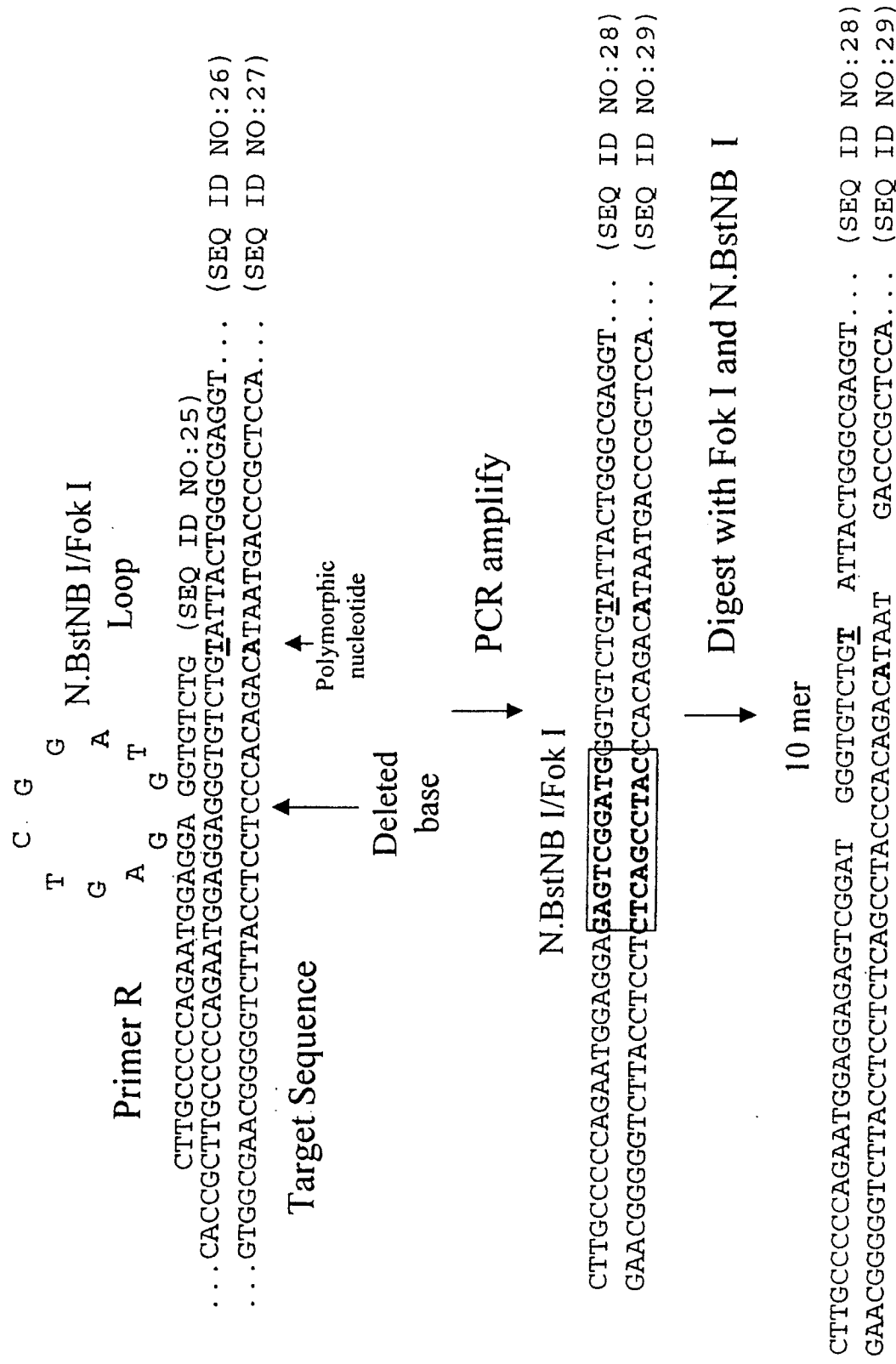


FIG. 9

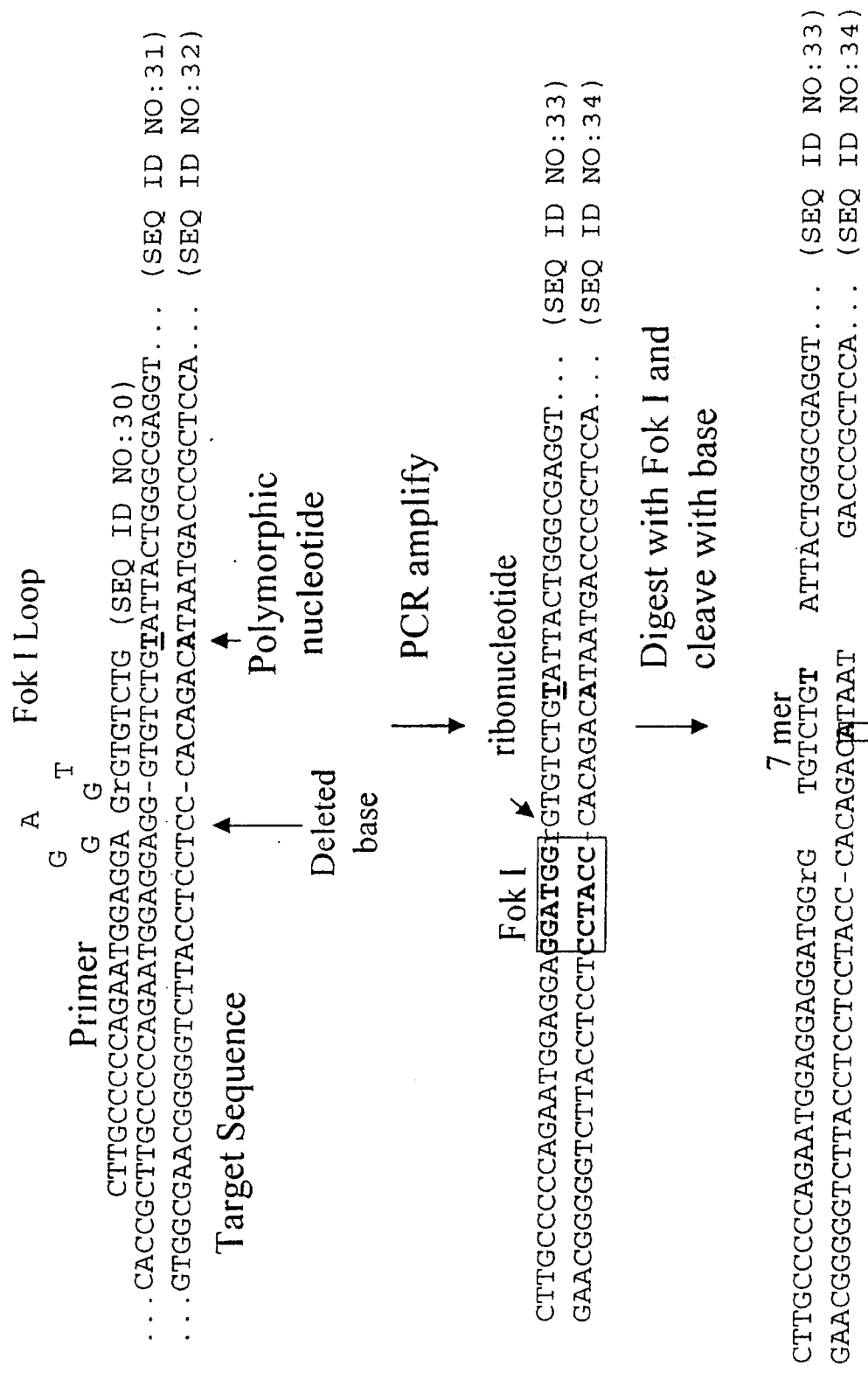


FIG. 10

METHODS FOR HAPLOTYPEING BASED ON PHYSICAL ALLELE SEPARATION

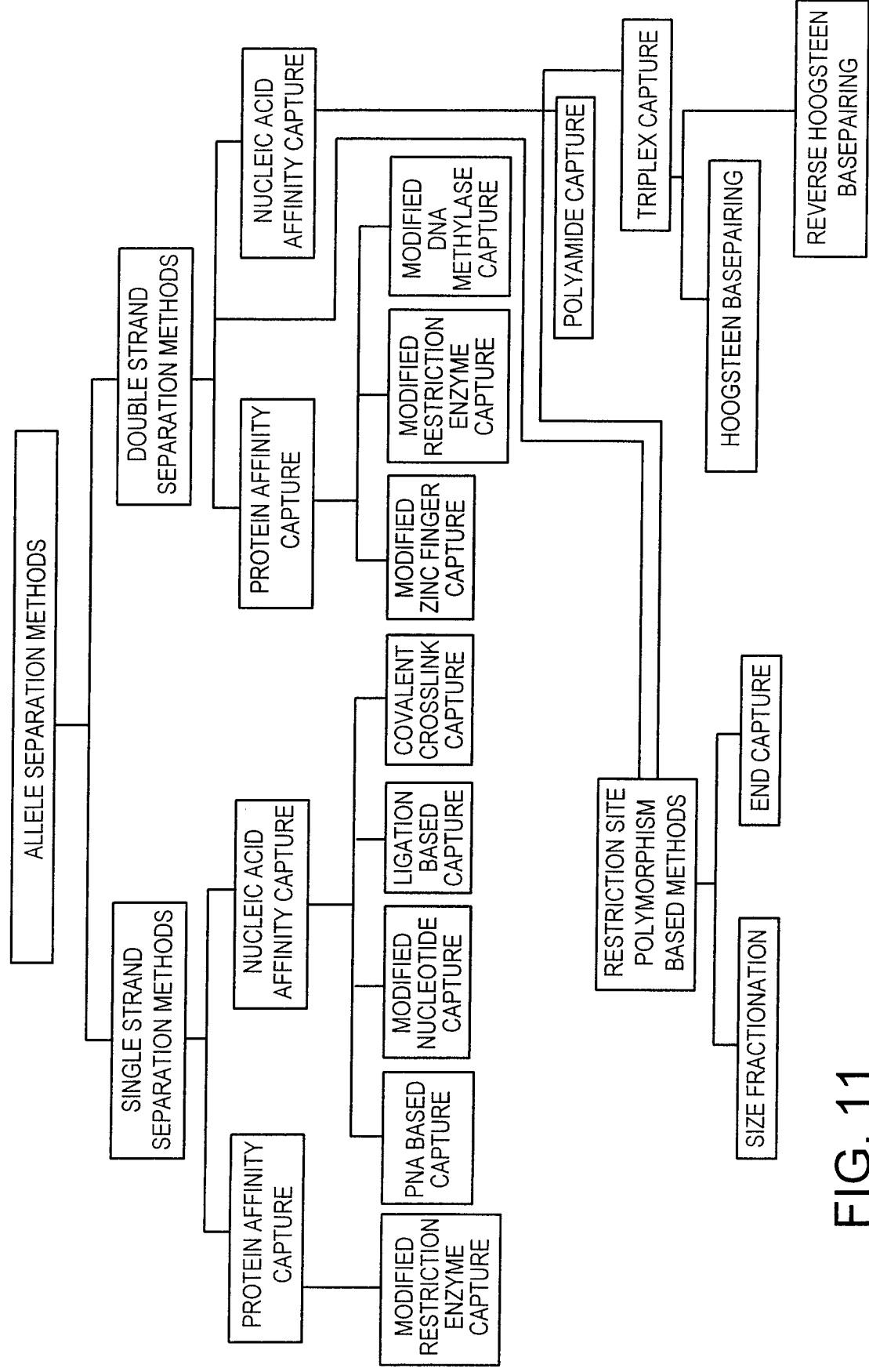
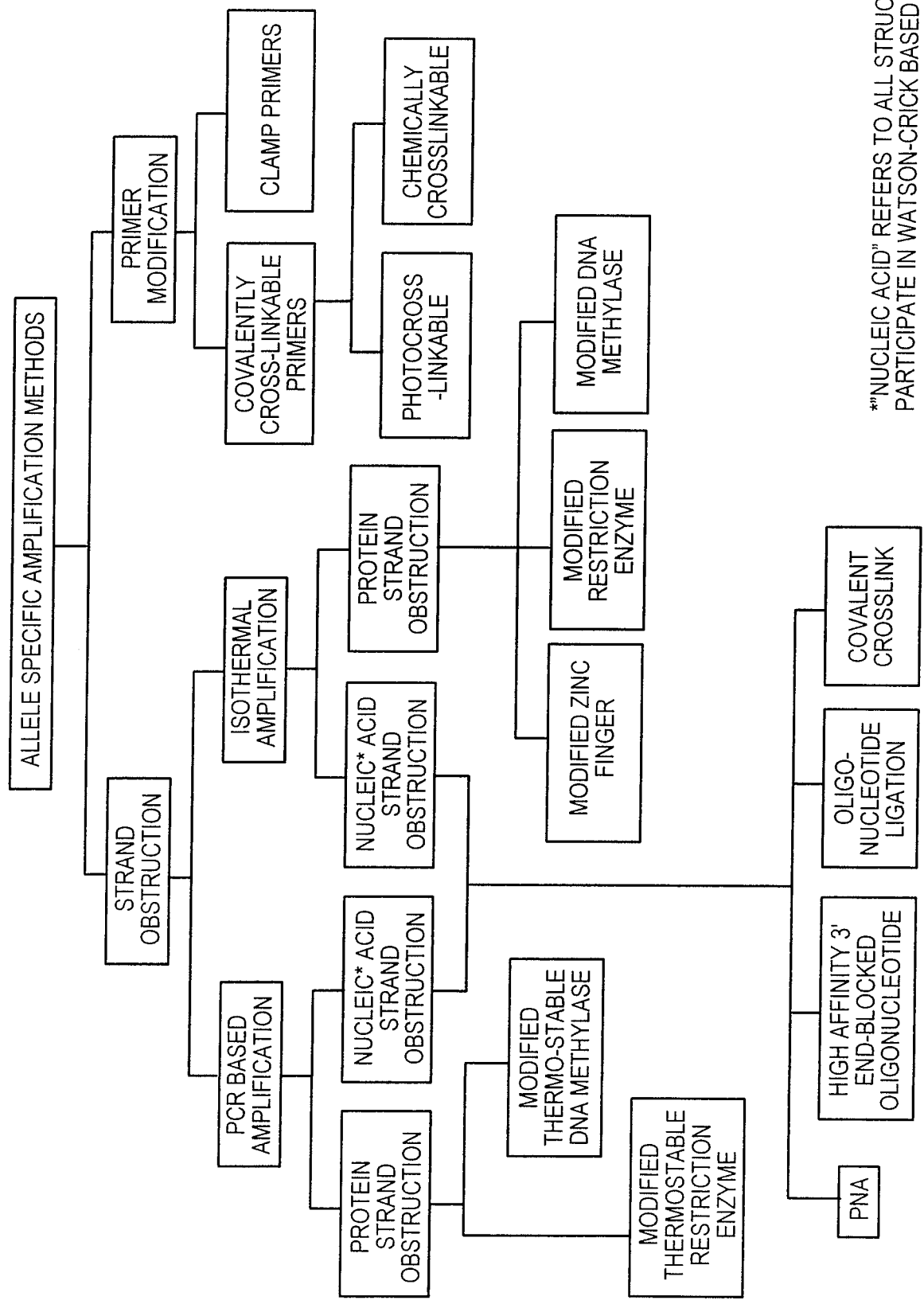


FIG. 11

METHODS FOR HAPLOTYPE ALLELE SPECIFIC AMPLIFICATION



*"NUCLEIC ACID" REFERS TO ALL STRUCTURES THAT PARTICIPATE IN WATSON-CRICK BASED PAIRING

FIG. 12

METHODS FOR HAPLOTYPING BASED ON ALLELE SPECIFIC RESTRICTION

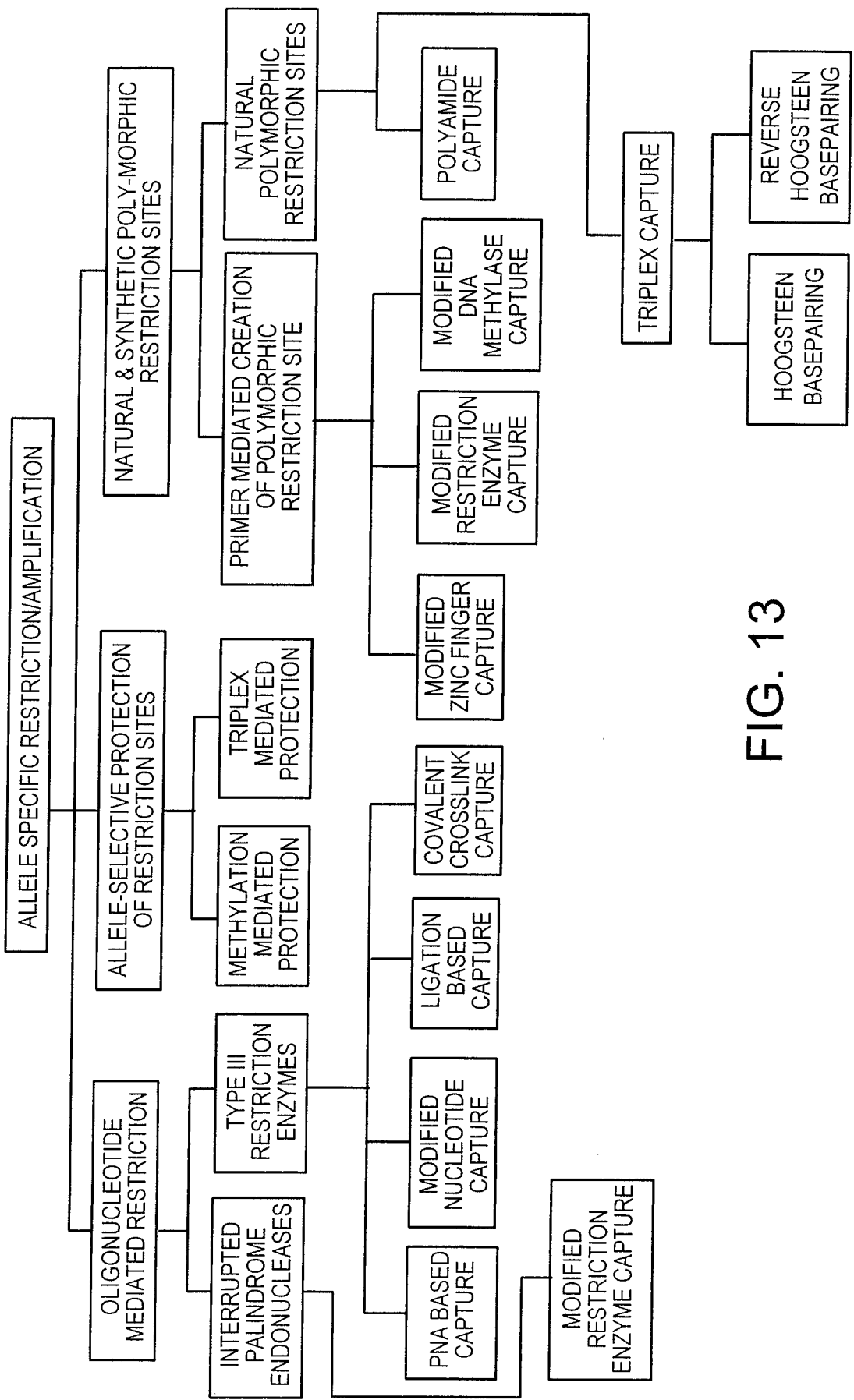


FIG. 13